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**Wober**

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(54) **NANOWIRE ARRAYS COMPRISING  
FLUORESCENT NANOWIRES AND  
SUBSTRATE**

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C01B 3/0021; C01B 31/02; C01B 3/00;  
G01J 1/04; G01J 1/42; G01J 5/08; B82Y  
20/00

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See application file for complete search history.

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(\*) Notice: Subject to any disclaimer, the term of this  
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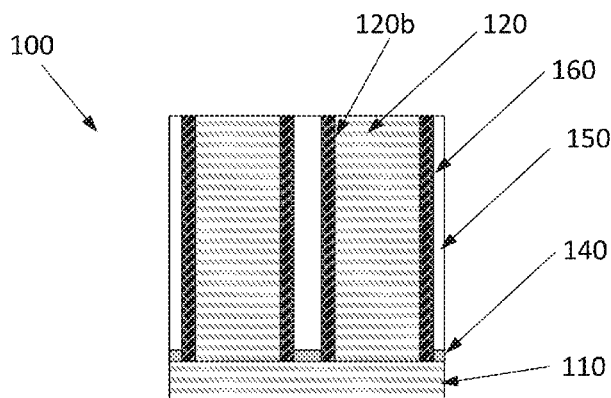
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(57) **ABSTRACT**

Described herein is a nanowire array, comprising a substrate,  
a plurality of fluorescent nanowires extending essentially per-  
pendicularly from the substrate and a reflective layer disposed  
on the substrate in areas between the fluorescent nanowires;  
wherein the fluorescent nanowires are operable to fluoresce at  
a wavelength of a collective mode of the nanowire array.

**23 Claims, 9 Drawing Sheets**



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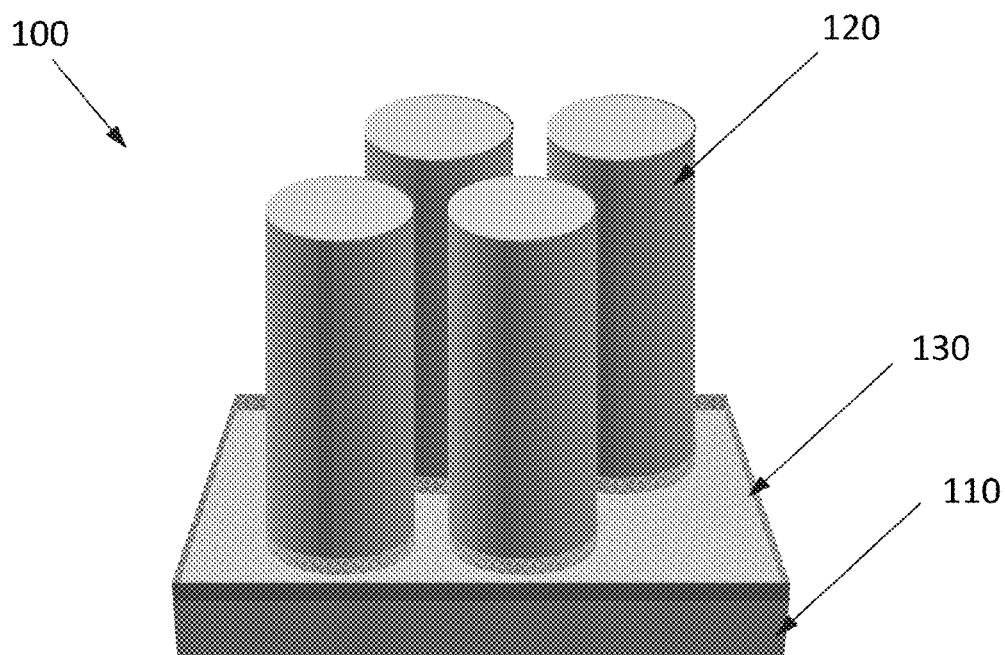


Fig. 1

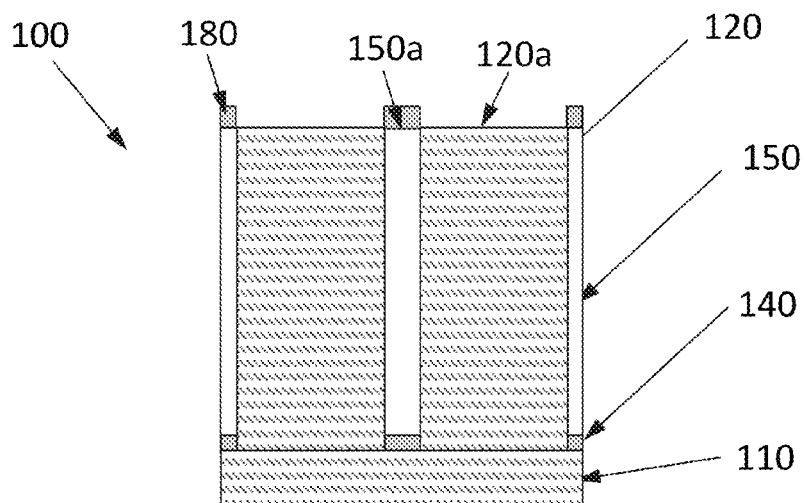


Fig. 2

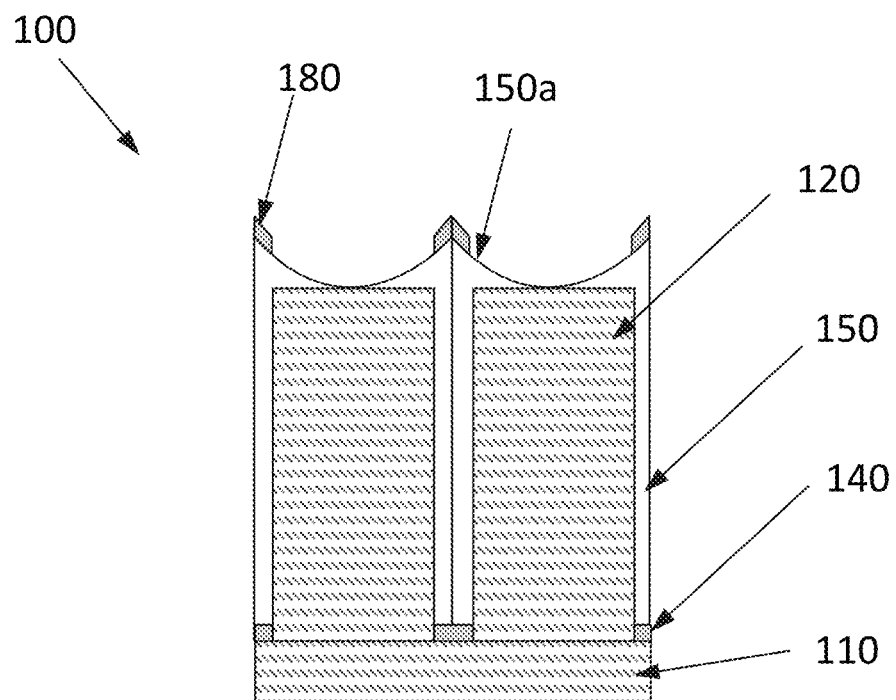


Fig. 3

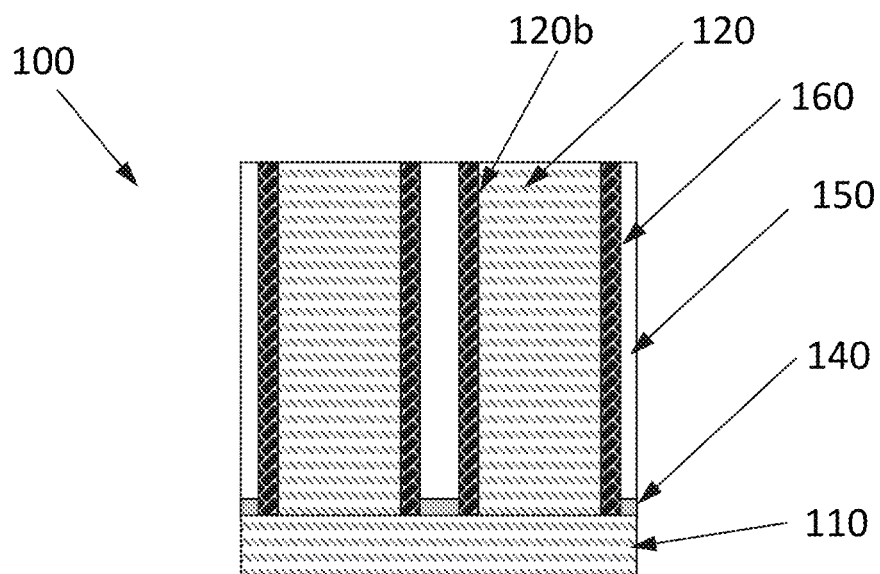


Fig. 4

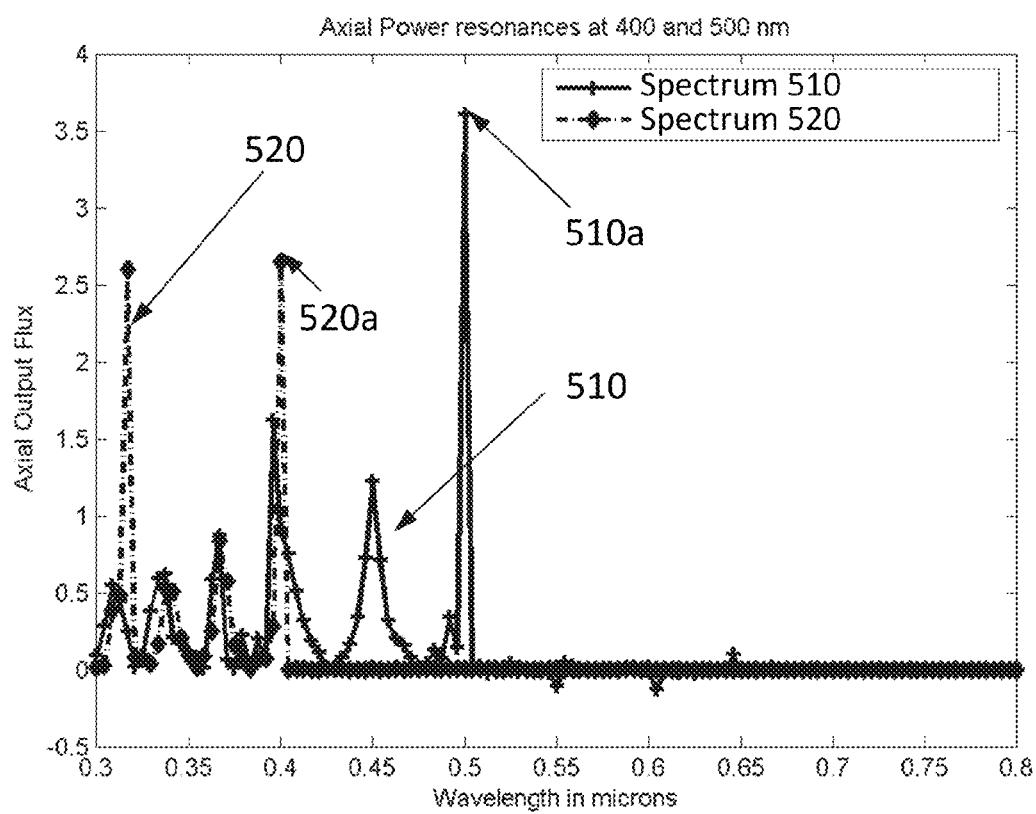


Fig. 5

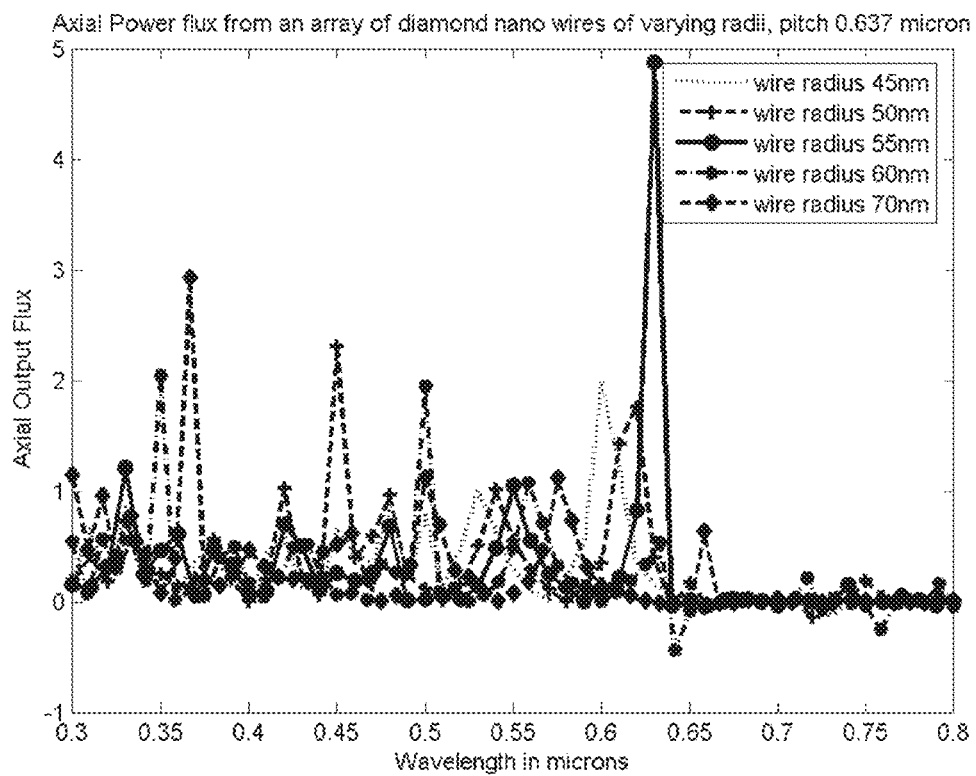


Fig. 6

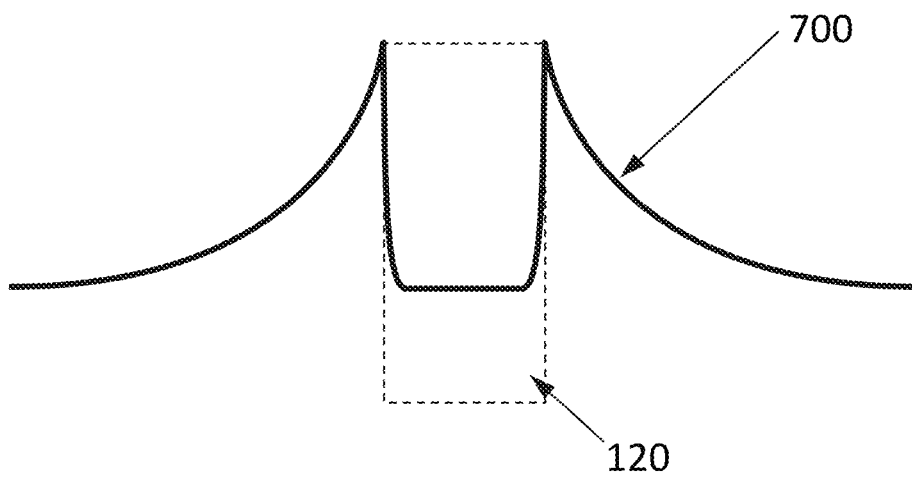


Fig. 7

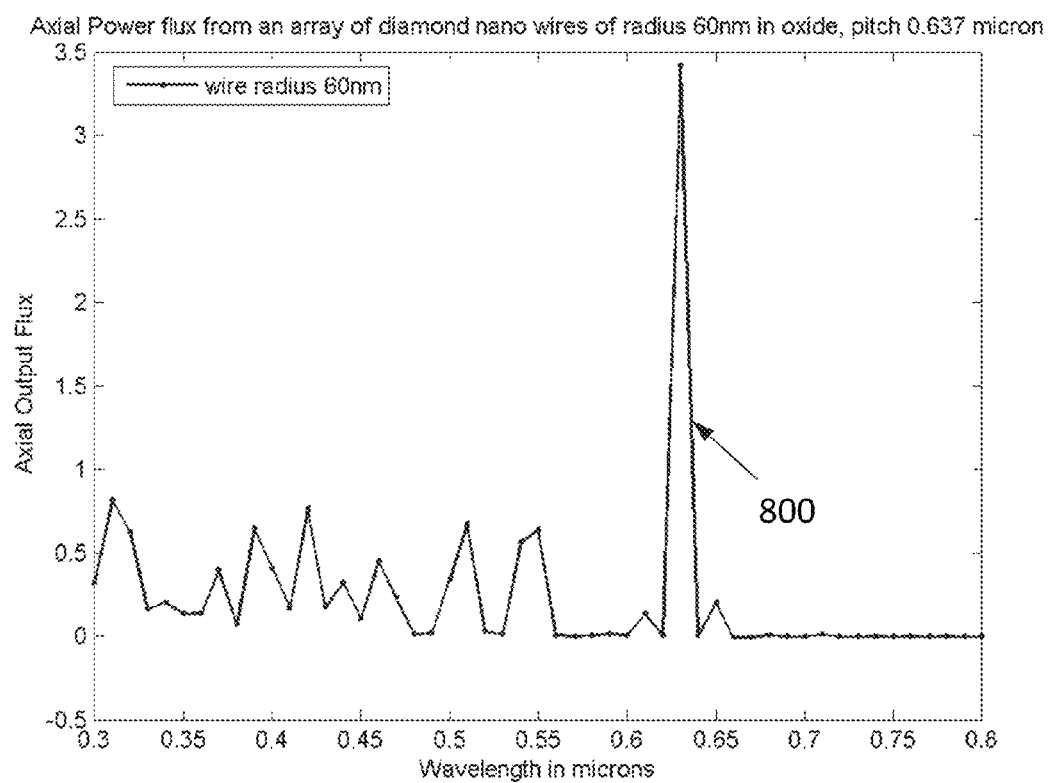


Fig. 8



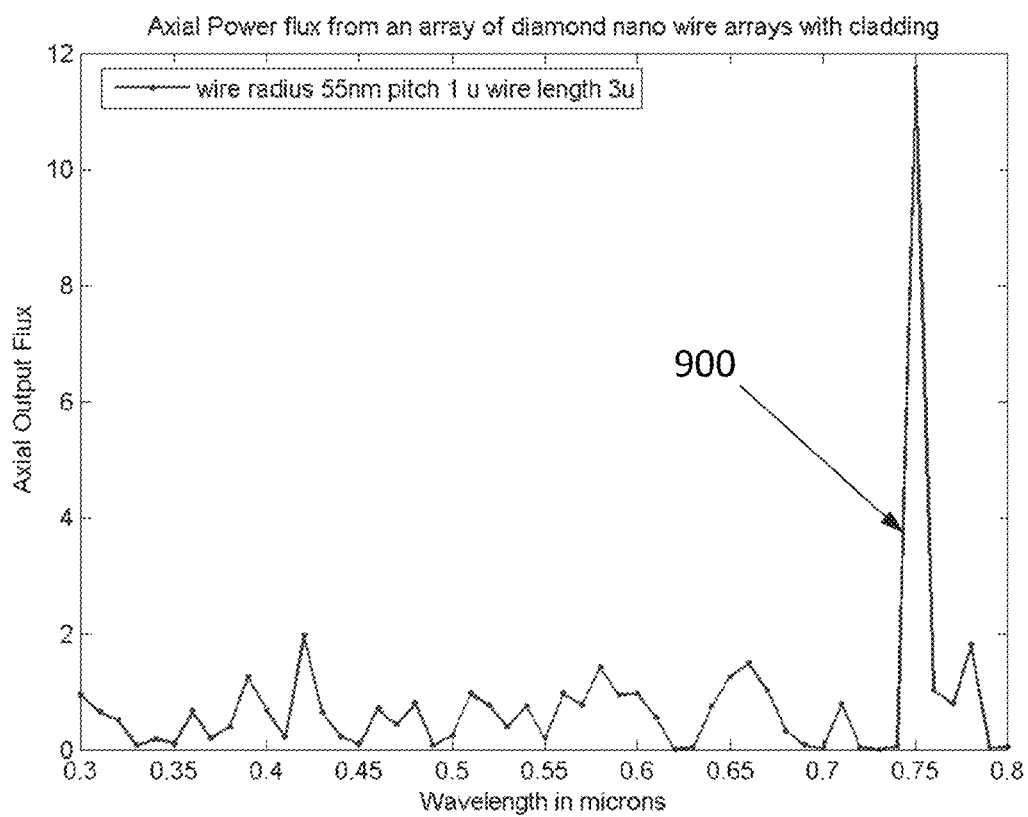


Fig. 9

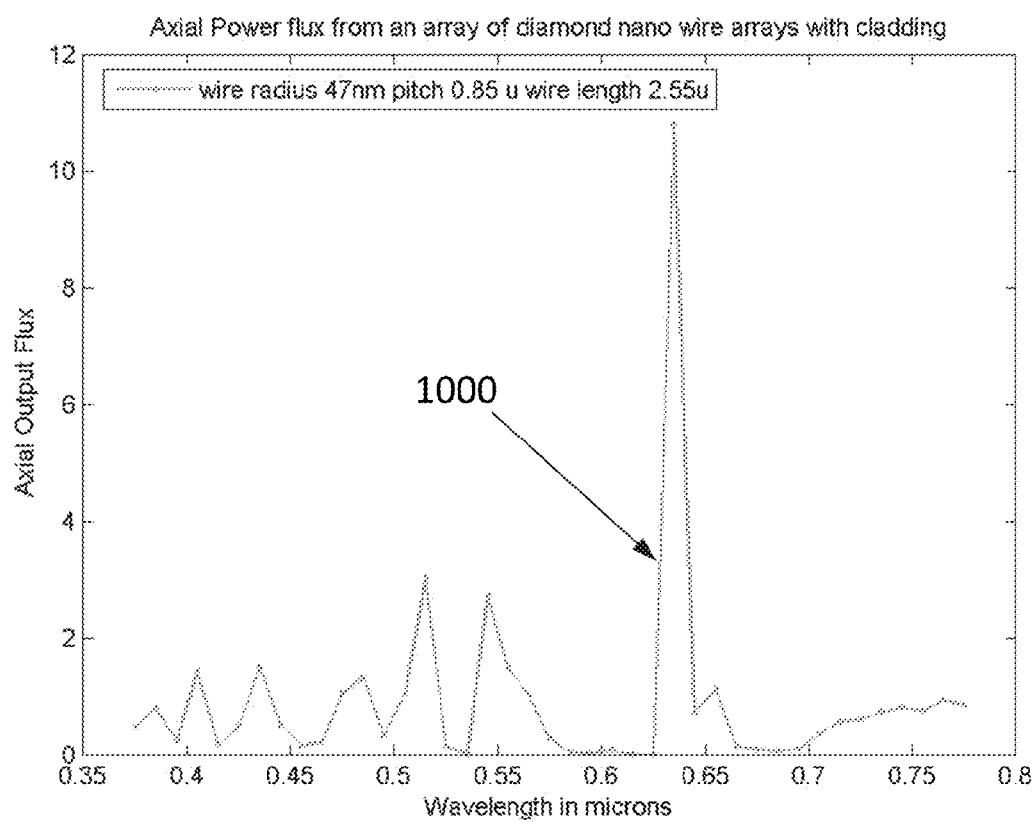


Fig. 10

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# NANOWIRE ARRAYS COMPRISING FLUORESCENT NANOWIRES AND SUBSTRATE

## CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. application Ser. No. 12/966,573, filed Dec. 13, 2013. This application is related to U.S. patent application Ser. No. 12/204,686, filed Sep. 4, 2008 (now U.S. Pat. No. 7,646,943, issued Jan. 12, 2010), Ser. No. 12/648,942, filed Dec. 29, 2009 (now U.S. Pat. No. 8,229,255, issued Jul. 24, 2012), Ser. No. 13/556,041, filed Jul. 23, 2012, Ser. No. 12/270,233, filed Nov. 13, 2008 (now U.S. Pat. No. 8,274,039, issued Sep. 25, 2012), Ser. No. 13/925,429, filed Jun. 24, 2013, Ser. No. 13/570,027, filed Aug. 8, 2012 (now U.S. Pat. No. 8,471,190, issued Jun. 25, 2013), Ser. No. 12/472,264, filed May 26, 2009 (now U.S. Pat. No. 8,269,985, issued Sep. 18, 2012), Ser. No. 13/621,607, filed Sep. 17, 2012 (now U.S. Pat. No. 8,514,411, issued Aug. 20, 2013), Ser. No. 13/971,523, filed Aug. 20, 2013 (now U.S. Pat. No. 8,810,808, issued Aug. 19, 2014), Ser. No. 12/472,271, filed May 26, 2009 (now abandoned), Ser. No. 12/478,598, filed Jun. 4, 2009 (now U.S. Pat. No. 8,546,742, issued Oct. 1, 2013), Ser. No. 14/021,672, filed Sep. 9, 2013, Ser. No. 12/573,582, filed Oct. 5, 2009 (now U.S. Pat. No. 8,791,470, issued Jul. 29, 2014), Ser. No. 14/274,448, filed May 9, 2014, Ser. No. 12/575,221, filed Oct. 7, 2009 (now U.S. Pat. No. 8,384,007, issued Feb. 26, 2013), Ser. No. 12/633,323, filed Dec. 8, 2009 (now U.S. Pat. No. 8,735,797, issued May 27, 2014), Ser. No. 14/068,864, filed Oct. 31, 2013, Ser. No. 14/281,108, filed May 19, 2014, Ser. No. 13/494,661, filed Jun. 12, 2012 (now U.S. Pat. No. 8,754,359, issued Jun. 17, 2014), Ser. No. 12/633,318, filed Dec. 8, 2009 (now U.S. Pat. No. 8,519,379, issued Aug. 27, 2013), Ser. No. 13/975,553, filed Aug. 26, 2013 (now U.S. Pat. No. 8,710,488, issued Apr. 29, 2014), Ser. No. 12/633,313, filed Dec. 8, 2009, Ser. No. 12/633,305, filed Dec. 8, 2009 (now U.S. Pat. No. 8,299,472, issued Oct. 30, 2012), Ser. No. 13/543,556, filed Jul. 6, 2012 (now U.S. Pat. No. 8,766,272, issued Jul. 1, 2014), Ser. No. 14/293,164, filed Jun. 2, 2014, Ser. No. 12/621,497, filed Nov. 19, 2009 (now abandoned), Ser. No. 12/633,297, filed Dec. 8, 2009 (now allowed), Ser. No. 12/982,269, filed Dec. 30, 2010, Ser. No. 12/967,880, filed Dec. 14, 2010 (now U.S. Pat. No. 8,748,799, issued Jun. 10, 2014), Ser. No. 14/291,888, filed May 30, 2014 12/966,514, filed Dec. 13, 2010, Ser. No. 12/974,499, filed Dec. 21, 2010 (now U.S. Pat. No. 8,507,840, issued Aug. 13, 2013), Ser. No. 12/966,535, filed Dec. 13, 2010, Ser. No. 12/910,664, filed Oct. 22, 2010, Ser. No. 12/945,492, filed Nov. 12, 2010, Ser. No. 13/047,392, filed Mar. 14, 2011 (now allowed), Ser. No. 14/450,812, filed Aug. 4, 2014, Ser. No. 13/048,635, filed Mar. 15, 2011 (now allowed), Ser. No. 13/106,851, filed May 12, 2011, Ser. No. 13/288,131, filed Nov. 3, 2011, Ser. No. 14/334,848, filed Jul. 18, 2014, Ser. No. 14/032,166, filed Sep. 19, 2013, Ser. No. 13/543,307, filed Jul. 6, 2012, Ser. No. 13/963,847, filed Aug. 9, 2013, Ser. No. 13/693,207, filed Dec. 4, 2012, 61/869,727, filed Aug. 25, 2013, Ser. No. 14/322,503, filed Jul. 2, 2014, and Ser. No. 14/311,954, filed Jun. 23, 2014, are each hereby incorporated by reference in their entirety.

## BACKGROUND

Nanostructures often exhibit fascinating physical properties not present in their bulk counterparts. Optical properties of nanostructures have been one of the recent research

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focuses. Tuning optical properties of nanostructures would facilitate their applications in the semiconductor, optics, and consumer electronics industry. In one example, optical properties of nanostructures can be controlled by their chemical composition. Chemical doping can change electronic structures of the materials semiconductor nanostructures are composed of, which in turn changes their interaction with light. In another example, arranging nanostructures into a regular lattice can yield optical properties individual nanostructures lack.

## SUMMARY

According to an embodiment, a nanowire array comprises a substrate, a plurality of fluorescent nanowires extending essentially perpendicularly from the substrate and a reflective layer at least disposed on the substrate in at least a portion of areas between the fluorescent nanowires; wherein the fluorescent nanowires are operable to fluoresce at a wavelength of a collective mode of the nanowire array. An electrically conductive material can be a material with essentially zero band gap. The electrical conductivity of an electrically conductive material is generally above  $10^3$  S/cm. A semiconductor can be a material with a finite band gap up to about 3 eV and general has an electrical conductivity in the range of  $10^3$  to  $10^{-8}$  S/cm. An electrically insulating material can be a material with a band gap greater than about 3 eV and generally has an electrical conductivity below  $10^{-8}$  S/cm. The term “fluorescent nanowires extending essentially perpendicular to the substrate” as used herein means that angles between the fluorescent nanowires and the substrate are from about  $85^\circ$  to  $90^\circ$ . The term “array” as used herein means a spatial arrangement having a particular order. The term “nanowire” as used herein means a structure that has a size constrained to at most 1000 nm in two dimensions and unconstrained in the other dimension. The term “disposed” as used herein means position, deposited, plated, or coated. The term “areas between the fluorescent nanowires” as used herein means areas on the substrate that is not covered by the fluorescent nanowires, as shown in FIGS. 2 and 3. The term “fluorescent nanowires” as used herein means nanowires that can fluoresce. The term “fluoresce” as used herein means to exhibit or undergo the phenomenon of fluorescence. The term “fluorescence” as used herein means the emission of light by a substance that has absorbed light or other electromagnetic radiation of a different wavelength. The term “fluoresced light” as used herein means emitted light from fluorescence of a substance. In most cases, emitted light has a longer wavelength, and therefore lower energy, than the absorbed radiation. However, when the absorbed electromagnetic radiation is intense, it is possible for one electron to absorb two photons; this two-photon absorption can lead to emission of radiation having a shorter wavelength than the absorbed radiation. Fluorescence has many practical applications, including mineralogy, gemology, chemical sensors, fluorescent labeling, dyes, biological detectors, and, most commonly, fluorescent lamps. For example, the common fluorescent lamp relies on fluorescence. Inside the glass tube is a partial vacuum and a small amount of mercury. An electric discharge in the tube causes the mercury atoms to emit ultraviolet light. The tube is lined with a coating of a fluorescent material, called the phosphor, which absorbs the ultraviolet and re-emits visible light.

According to an embodiment, the substrate comprises a material comprising silicon, sapphire, glass silicon oxide, diamond, ZnO, a transparent non-absorbing dielectric material or a combination thereof. Glass is a non-crystalline solid material. The most prevalent type of glass, used for centuries

in windows and drinking vessels, is soda-lime glass, made of about 75% silica ( $\text{SiO}_2$ ) plus  $\text{Na}_2\text{O}$ ,  $\text{CaO}$ , and several minor additives. Often, the term glass is used in a restricted sense to refer to this specific use. In science, however, the term glass is usually defined in a much wider sense, including every solid that possesses a non-crystalline (i.e. amorphous) structure and that exhibits a glass transition when heated towards the liquid state. In this wider sense, glasses can be made of quite different classes of materials: metallic alloys, ionic melts, aqueous solutions, molecular liquids, and polymers. Of these, polymer glasses (acrylic glass, polyethylene terephthalate) are the most important; for many applications (bottles, eye-wear) they are a lighter alternative to traditional silica glasses.

According to an embodiment, the substrate has a thickness of about 0.1 to 1 mm.

According to an embodiment, the substrate comprises substantially a same material as or different materials from the fluorescent nanowires.

According to an embodiment, the reflective layer comprises a material selected from a group consisting of  $\text{ZnO}$ ,  $\text{Al}$ ,  $\text{Au}$ ,  $\text{Ag}$ ,  $\text{Pd}$ ,  $\text{Cr}$ ,  $\text{Cu}$ , and a combination thereof.

According to an embodiment, the reflective layer has a reflectance of at least 50% for fluoresced light from the fluorescent nanowires. The term "reflectance" as used herein means the fraction of incident electromagnetic power that is reflected at an interface.

According to an embodiment, the reflective layer has a thickness of at least 5 nm.

According to an embodiment, the reflective layer is also disposed on the substrate in at least a portion of areas covered by the fluorescent nanowires.

According to an embodiment, the fluorescent nanowires are at least partially embedded in a material. The term "embed" as used herein means to surround or cover something closely.

According to an embodiment, the material comprises silicon oxide, silicon nitride,  $\text{ZnO}$ ,  $\text{Al}_2\text{O}_3$ , diamond or a combination thereof. The material can also comprise any suitable transparent and low absorption dielectric material.

According to an embodiment, a surface the material is substantially coextensive with a surface of each of the fluorescent nanowires such the surface of each of the fluorescent nanowires is exposed. The term "exposed" as used herein means not embedded by the material.

According to an embodiment, the fluorescent nanowires are entirely embedded in the material without any exposed surface.

According to an embodiment, a surface of the material has a concave shape configured to focus fluoresced light from the fluorescent nanowires.

According to an embodiment, a refractive index of the material is smaller than a refractive index of the fluorescent nanowires.

According to an embodiment, the nanowire array further comprises a partial reflective layer deposited on a surface of the material in at least a portion of areas around the surface of each of the fluorescent nanowires wherein no portion of the surface of each of the fluorescent nanowires overlaps the partial reflective layer when viewed from a direction perpendicular to the substrate.

According to an embodiment, the fluorescent nanowires comprise one or more fluorescent material. The term "fluorescent material" as used herein means a material that can fluoresce.

According to an embodiment, the fluorescent nanowires comprise a material selected from the group consisting of inorganic fluorescent materials, organic fluorescent materials

and a combination thereof. Organic fluorescent materials, often referred to as organic dyes, include a large number of organic molecules that can fluoresce. Several examples of organic fluorescent materials are acridine orange, SYBR Green (N',N'-dimethyl-N-[4-[(E)-(3-methyl-1,3-benzothiazol-2-ylidene)methyl]-1-phenylquinolin-1-ium-2-yl]-N-propylpropane-1,3-diamine), cyanines, DiOC<sub>6</sub> (3-Hexyl-2-[3-(3-hexyl-2(3H)benzoxazolylidene)-1-propenyl]benzoxazolium iodide, CAS number 53213-82-4), Nile blue, Nile red, oxazin dyes, phenanthridine dyes, Rhodamine dyes, etc. Inorganic fluorescent materials include a large number of inorganic materials that can fluoresce. Several examples of inorganic fluorescent materials are  $\text{Zn}_2\text{SiO}_4\text{:Mn}$ ,  $\text{GaN}$ ,  $\text{ZnO:Zn}$ ,  $\text{ZnS:Ag}$ ,  $\text{CdS:Ag}$ ,  $\text{MgF}_2\text{:Mn}$ ,  $\text{InBO}_3\text{:Tb}$ ,  $\text{LaPO}_4\text{:Ce}$ , etc. The inorganic fluorescent materials can also be suitable quantum dots (e.g. silicon). A quantum dot is a semiconductor whose excitons are confined in all three spatial dimensions.

According to an embodiment, the fluorescent nanowires are one or more non-fluorescent material with one or more fluorescent material embedded therein and covered thereby. The term "non-fluorescent material" as used herein means a material that cannot fluoresce.

According to an embodiment, the fluorescent nanowires are diamond with color centers embedded therein. Diamond is an allotrope of carbon, where the carbon atoms are arranged in a variation of the face-centered cubic crystal structure called a diamond lattice. The term "color center" as used herein means a point lattice defect which produces optical absorption bands in an otherwise transparent crystal. such as the alkali halides, alkaline earth fluorides, or metal oxides. Color centers are found in a wide range of materials. Color centers can be produced by gamma radiation or x-radiation, by addition of impurities or excess constituents, and sometimes through electrolysis. A well-known example is that of the F-center in alkali halides such as sodium chloride,  $\text{NaCl}$ . The designation F-center comes from the German word Farbe, which means color. F-centers in  $\text{NaCl}$  produce a band of optical absorption toward the blue end of the visible spectrum; thus the colored crystal appears yellow under transmitted light. On the other hand,  $\text{KCl}$  with F-centers appears magenta, and  $\text{KBr}$  appears blue.

According to an embodiment, the color centers are nitrogen vacancy defects.

According to an embodiment, the color centers are embedded randomly in the fluorescent nanowires.

According to an embodiment, at least 10% of the fluorescent nanowires have at least one color center.

According to an embodiment, the nanowire array further comprises a cladding layer surrounding a sidewall of each of the fluorescent nanowires wherein the cladding layer is a material substantially transparent to fluoresced light from the fluorescent nanowires. The term "cladding layer" as used herein means a layer of substance surrounding the structures.

According to an embodiment, a refractive index of cladding layer is smaller than a refractive index of the fluorescent nanowires.

According to an embodiment, the fluorescent nanowires are embedded in a material and wherein a refractive index of cladding layer is greater than the refractive index of the material.

According to an embodiment, the fluorescent nanowires have a substantially uniform length. The term "substantially uniform length" as used herein means all lengths of the fluorescent nanowires are from 90% to 110% of a mean value of all the lengths.

According to an embodiment, fluoresced light of the fluorescent nanowires is substantially spatially and/or temporally

coherent. Coherence describes all properties of the correlation between physical quantities of a wave. When interfering, two waves can add together to create a larger wave (constructive interference) or subtract from each other to create a smaller wave (destructive interference), depending on their relative phase. Two waves are said to be coherent if they have a constant relative phase. The degree of coherence is measured by the interference visibility, a measure of how perfectly the waves can cancel due to destructive interference. Temporal coherence is the measure of the average correlation between the value of a wave at any pair of times, separated by delay  $T$ . Temporal coherence tells us how monochromatic a source is. In other words, it characterizes how well a wave can interfere with itself at a different time. Spatial coherence describes the ability for two points in space,  $x_1$  and  $x_2$ , in the extent of a wave to interfere, when averaged over time. More precisely, the spatial coherence is the cross-correlation between two points in a wave for all times.

According to an embodiment, the wavelength of the collective mode of the nanowire array is substantially equal to a pitch of the nanowire array, which as used herein means the wavelength of the collective mode of the nanowire array is from 90% to 110% of the pitch of the nanowire array.

According to an embodiment, a length of the fluorescent nanowires is a multiple of an effective wavelength  $\lambda_{eff}$  defined as  $\lambda_{eff} = \lambda_{air} / n_{eff}$ , wherein  $\lambda_{air}$  is the wavelength of interest and  $n_{eff}$  is an effective refractive index of the nanowire array at  $\lambda_{air}$ .

According to an embodiment, the collective mode is an air mode.

According to an embodiment, a pitch of the nanowire array is about 100 nm to 10 micron, a length of the fluorescent nanowires is from about 100 nm to 10 microns, and/or a radius of the fluorescent nanowires is about 20 to 200 nm.

According to an embodiment, a light source comprises one or more of the nanowire array.

According to an embodiment, a method of using the nanowire array comprises exposing the nanowire array to light.

According to an embodiment, the light has a shorter wavelength than the wavelength of the collective mode of the nanowire array.

According to an embodiment, the light is provided by a source selected from the group consisting of LEDs, fluorescent lamps, mercury-vapor lamps, sodium-vapor lamps, discharge lamps, sunlight, incandescent lamps and/or laser.

According to an embodiment, a method of designing a nanowire array that fluoresces at a wavelength of interest comprises: selecting the wavelength of interest; selecting a pitch of the nanowire array to be substantially equal to the wavelength of interest; selecting a radius of fluorescent nanowires of the nanowire array that maximizes fluorescence intensity of the nanowire array at the wavelength of interest; selecting a material of the fluorescent nanowires such that the fluorescent nanowires fluoresce at the wavelength of interest. The term "fluorescence intensity" as used herein means light intensity of fluoresced light.

According to an embodiment, a method of manufacturing the nanowire array comprises providing a substrate; growing fluorescent nanowires on the substrate such that the fluorescent nanowires extend essentially perpendicularly from the substrate and the fluorescent nanowires are operable to fluoresce at a wavelength of a collective mode of the nanowire array; depositing the reflective layer at least on the substrate in at least a portion of areas between the fluorescent nanowires. A resist layer as used herein means a thin layer used to transfer a pattern to the substrate which the resist layer is

deposited upon. A resist layer can be patterned via lithography to form a (sub)micrometer-scale, temporary mask that protects selected areas of the underlying substrate during subsequent processing steps. The resist is generally proprietary mixtures of a polymer or its precursor and other small molecules (e.g. photoacid generators) that have been specially formulated for a given lithography technology. Resists used during photolithography are called photoresists. Resists used during e-beam lithography are called e-beam resists. A lithography technique can be photolithography, e-beam lithography, holographic lithography. Photolithography is a process used in microfabrication to selectively remove parts of a thin film or the bulk of a substrate. It uses light to transfer a geometric pattern from a photo mask to a light-sensitive chemical photo resist, or simply "resist," on the substrate. A series of chemical treatments then engraves the exposure pattern into the material underneath the photo resist. In complex integrated circuits, for example a modern CMOS, a wafer will go through the photolithographic cycle up to 50 times. E-beam lithography is the practice of scanning a beam of electrons in a patterned fashion across a surface covered with a film (called the resist), ("exposing" the resist) and of selectively removing either exposed or non-exposed regions of the resist ("developing"). The purpose, as with photolithography, is to create very small structures in the resist that can subsequently be transferred to the substrate material, often by etching. It was developed for manufacturing integrated circuits, and is also used for creating nanotechnology artifacts.

According to an embodiment, a method of manufacturing the nanowire array of comprises: providing a substrate having a layer of fluorescent material; coating a resist layer on the layer of fluorescent material; patterning the resist layer using a lithographic technique such that a pattern corresponding to a plurality of fluorescent nanowires is formed in the resist layer; forming the fluorescent nanowires by etching the layer of fluorescent material such that the fluorescent nanowires extend essentially perpendicularly from the substrate and the fluorescent nanowires are operable to fluoresce at a wavelength of a collective mode of the nanowire array; depositing a reflective layer at least on the substrate in at least a portion of areas between the fluorescent nanowires.

According to an embodiment, the method further comprises a step of coating a cladding layer surrounding a sidewall of the fluorescent nanowires; and/or a step of filling space between the fluorescent nanowires with a material.

According to an embodiment, a method of manufacturing a nanowire array comprising: providing a substrate; growing nanowires on the substrate such that the nanowires extend essentially perpendicularly from the substrate; depositing a fluorescent material on or embedding a fluorescent material in the nanowires followed by an optional annealing step such that the nanowires are operable to fluoresce at a wavelength of a collective mode of the nanowire array; depositing the reflective layer at least on the substrate in at least a portion of areas between the nanowires.

According to an embodiment, a method of manufacturing a nanowire array comprising: providing a substrate having a layer of a material; coating a resist layer on the layer of the material; patterning the resist layer using a lithographic technique such that a pattern corresponding to a plurality of nanowires is formed in the resist layer; forming the nanowires by etching the layer of fluorescent material such that the nanowires extend essentially perpendicularly from the substrate; depositing a fluorescent material on or embedding a fluorescent material in the nanowires followed by an optional annealing step such that the nanowires are operable to fluo-

resce at a wavelength of a collective mode of the nanowire array; depositing a reflective layer at least on the substrate in at least a portion of areas between the nanowires.

According to an embodiment, the method further comprises a step of coating a cladding layer surrounding a side-wall of the nanowires; and/or a step of filling space between the nanowires with a material.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Embodiments of the present disclosure will now be disclosed, by way of example only, with reference to the accompanying schematic drawings in which corresponding reference symbols indicate corresponding parts, in which:

FIG. 1 shows a schematic perspective view of a nanowire array, according to an embodiment.

FIG. 2 shows a schematic cross-sectional view of the nanowire array of FIG. 1, according to an embodiment.

FIG. 3 shows an alternative schematic cross-sectional view of the nanowire array of FIG. 1, according to an embodiment.

FIG. 4 shows yet another schematic cross-sectional view of the nanowire array of FIG. 1, according to an embodiment.

FIG. 5 shows simulated fluorescence spectra of two exemplary nanowire arrays in axial directions thereof.

FIG. 6 shows simulated fluorescence spectra of several exemplary nanowire arrays in axial directions thereof, all of which have a same pitch of 637 nm and different radii.

FIG. 7 shows a schematic light distribution of an air mode of the nanowire array.

FIG. 8 shows a simulated fluorescence spectrum of an exemplary nanowire array in axial directions thereof, wherein the fluorescent nanowires are embedded in oxide.

FIG. 9 shows a simulated fluorescence spectrum of an exemplary nanowire array in axial directions thereof, wherein the fluorescent nanowires have a cladding layer.

FIG. 10 shows a simulated fluorescence spectrum of an exemplary nanowire array in axial directions thereof, wherein the fluorescent nanowires have a cladding layer and wherein the nanowire array of FIG. 10 has dimensions to scale with the nanowire array of FIG. 9 and a peak position at 637 nm.

#### DETAILED DESCRIPTION

In the following detailed description, reference is made to the accompanying drawings, which form a part thereof. In the drawings, similar symbols typically identify similar components, unless the context dictates otherwise. The illustrative embodiments described in the detail description, drawings, and claims are not meant to be limiting. Other embodiments may be utilized, and other changes may be made, without departing from the spirit or scope of the subject matter presented here.

FIGS. 1 and 2 are a schematic perspective view and a schematic cross-sectional view of a nanowire array 100, respectively, according to an embodiment. The nanowire array 100 comprises a substrate 110, a plurality of fluorescent nanowires 120, a reflective layer 140 disposed on the substrate 110 in areas between the fluorescent nanowires 120. The fluorescent nanowires 120 can be embedded in a material 150.

The substrate 110 can be any suitable material, such as, silicon, sapphire, glass, silicon oxide, etc. The substrate 110 and the fluorescent nanowires 120 can be the same material or different materials. The substrate 110 can have any suitable thickness such as 0.1-1 mm. Examples of the substrate 110 include a silicon wafer, a sapphire wafer, and a glass wafer.

The reflective layer 140 can be any suitable material, such as ZnO, Al, Au, Ag, Pd, Cr, Cu, a combination thereof, etc. The reflective layer 140 preferably has a reflectance of at least 50%, more preferably has a reflectance of at least 70%, most preferably has a reflectance of at least 90%, for fluoresced light from the fluorescent nanowires 120. The reflective layer 140 has a thickness of preferably at least 5 nm, more preferably at least 20 nm. In addition to the areas between the fluorescent nanowires 120, the reflective layer 140 can also be disposed on the substrate 110 in areas under the fluorescent nanowires 120.

The material 150 in which the are embedded can be any suitable material substantially transparent to fluoresced light from the fluorescent nanowires 120. For example, the material 150 can be silicon oxide, silicon nitride or a combination thereof. A surface 150a of the material 150 can be substantially coextensive with surfaces 120a of the fluorescent nanowires 120 such that the surfaces 120a of the fluorescent nanowires 120 are exposed. Alternatively, the fluorescent nanowires 120 can be entirely buried in the material 150 without any exposed surface. As shown in FIG. 3, the surface 150a of the material 150 can have a concave shape effective to focus fluoresced light from the fluorescent nanowires 120. A refractive index of the material 150 is preferably smaller than a refractive index of the fluorescent nanowires 120. A partial reflective layer 180 can be deposited on the surface 150a in areas around the surfaces 120a wherein no portion of the surfaces 120a overlaps the partial reflective layer 180. The term "partial reflective layer" as used herein means a layer with a reflectance less than 100%. For example, a partial reflective layer can be a layer of Ag or Al with a thickness less than 10 nm.

The fluorescent nanowires 120 can comprise one or more suitable fluorescent material, such as GaN, ZnO, organic fluorescent materials, or a combination thereof. The fluorescent nanowires 120 can also be one or more non-fluorescent material with one or more fluorescent material embedded therein and covered thereby, such as diamond with color centers embedded therein. Fluorescence is the emission of light by a substance that has absorbed light or other electromagnetic radiation of a different wavelength from emitted light. In most cases, the emitted light has a longer wavelength, and therefore lower energy, than the absorbed radiation. In an example, the fluorescent nanowires 120 are diamond with nitrogen vacancy defects as the color centers. The color centers can be embedded randomly in the fluorescent nanowires 120. In a given nanowire array 100, at least 100%, 75%, 50%, 25%, or 10% of the fluorescent nanowires 120 have at least one color center. A nitrogen vacancy defect (also referred to as a nitrogen vacancy center) is one of numerous point defects in diamond consisting of a substitutional nitrogen atom and a vacancy in an adjacent site of the diamond lattice. Diamond with embedded nitrogen vacancy defects can be created by irradiating nitrogen doped diamond with 1-2 MeV electrons followed by annealing in vacuum at about 850-1000° C. for about 2-20 hours. Nitrogen vacancy defects are resistant to photobleaching, i.e., fluorescence intensity of nitrogen vacancy defects remains unchanged after several hours of continuous laser irradiation in the saturation regime. The saturation regime as used here in means a range of laser irradiation intensity in which fluorescence intensity of the nitrogen vacancy defects is independent from the laser irradiation intensity. Nitrogen vacancy defects can have high quantum yield at temperatures up to in excess of 500 K. A nitrogen vacancy defect can fluoresce at wavelengths such as 637 nm and 531 nm, depending on the symmetry of the nitrogen atom of the nitrogen vacancy defect relative to the

diamond lattice. The term “quantum yield” gives the efficiency of the fluorescence process and is defined as the ratio of the number of photons emitted to the number of photons absorbed. The fluorescent nanowires **120** can have any cross-sectional shape. For example, the fluorescent nanowires **120** can be cylinders or prisms with elliptical, circular, rectangular, polygonal cross-sections.

As shown in FIG. 4, the fluorescent nanowires **120** can have a cladding layer **160** surrounding a sidewall **120b** of each of the fluorescent nanowires **120**. The cladding layer **160** can be any suitable material substantially transparent to fluoresced light from the fluorescent nanowires **120**. If the fluorescent nanowires **120** are embedded in the material **150**, a refractive index of cladding layer **160** preferably is greater than the refractive index of the material **150** and smaller than the refractive index of the fluorescent nanowires **120**.

The fluorescent nanowires **120** preferably are arranged in an array, such as a rectangular array, a hexagonal array, a square array, concentric ring. The term “pitch” is defined as a distance of a fluorescent nanowire **120** to a nearest neighbor of the fluorescent nanowire **120** along a direction parallel to the substrate **110**. The term “array” as used herein means a spatial arrangement having a particular order. The fluorescent nanowires **120** preferably have a substantially uniform length.

The pitch of the nanowire array **100** and the radius, length, material of the fluorescent nanowires **120**, and the material **150** are selected such that the fluorescent nanowires **120** fluoresce at a wavelength of a collective mode of the nanowire array **100**. The pitch of the nanowire array **100** is preferably 100 nm to 10 microns, more preferably 300 nm to 1 micron. The radius of the fluorescent nanowires **120** is preferably 20 to 200 nm, more preferably 30 to 80 nm. The length of the fluorescent nanowires **120** is preferably from 100 nm to 10 microns, more preferably from 800 nm to 5 microns. The collective mode is a mode of the entire nanowire array **100** instead of a mode of individual fluorescent nanowires **120**. When the fluorescent nanowires **120** fluoresce at the collective mode of the nanowire array **100**, fluoresced light of the fluorescent nanowires **120** is substantially spatially and/or temporally coherent. Fluoresced light from such a nanowire array **100** has a much higher intensity than fluoresced light from an individual fluorescent nanowire and remains coherent. Namely, the nanowire array **100** can be a coherent light source. A coherent light source is a light source whose emitted light is substantially coherent, temporally, spatially, or both. A coherent light source can have application such as in quantum communication, in quantum cryptography, and as laser. The term “mode” as used herein means a possible standing electromagnetic wave in the nanowire array **100**. A standing wave, also known as a stationary wave, is a wave that remains in a constant position. In a stationary medium, a standing wave can occur as a result of interference between two waves traveling in opposite directions. Not all standing waves are possible in a given system. Possible standing waves are dictated by characteristics of the system, such as, physical dimensions, shapes, chemical composition and/or characteristics of an environment in the vicinity of the system. The wavelength of the standing wave of a mode is called the wavelength of the mode.

FIG. 5 shows simulated fluorescence spectra of two exemplary nanowire arrays in axial directions thereof. The axial direction of a nanowire array means the direction perpendicular to the substrate of the nanowire array. The simulation is done using the finite difference time domain (FDTD) method. The FDTD method is a method of numerically simulating propagation of light in a structure and can be used to predict

detailed characteristics of the propagation. Both spectra **510** and **520** show sharp peaks such as the peaks **510a** and **520a**. These peaks **510a** and **520a** are at wavelengths of collective modes of the exemplary nanowire arrays, respectively. Spectrum **510** is a fluorescence spectra of one of the exemplary nanowire arrays wherein the fluorescent nanowires are diamond and have a radius of 50 nm and pitch of 500 nm. Spectrum **520** is a fluorescence spectra of the other one of the exemplary nanowire arrays wherein the fluorescent nanowires are diamond, have a radius of 60 nm and pitch of 400 nm. In both spectra **510** and **520**, the highest peaks **510a** and **520a** are at wavelengths substantially equal to the pitches of the exemplary nanowire arrays, respectively. This observation leads to a method of designing a nanowire array that fluoresces at a wavelength of interest, the method comprising: selecting the wavelength of interest; selecting a pitch of the nanowire array to be substantially equal to the wavelength of interest; selecting a radius of fluorescent nanowires of the nanowire array that maximizes fluorescence intensity of the nanowire array at the wavelength of interest; selecting a material of the fluorescent nanowires such that the fluorescent nanowires fluoresce at the wavelength of interest.

FIG. 6 shows results of this method of design for a wavelength of interest at about 630 nm. Among five nanowire arrays all with a pitch of 637 nm but different radii of fluorescent nanowires, one nanowire arrays with fluorescent nanowires with a radius of about 55 nm maximizes fluorescence intensity of the nanowire array at the wavelength of interest.

Once a pitch of a nanowire array, a radius and a material of fluorescent nanowires in the nanowire array are determined, for examples, by the methods above, length of the fluorescent nanowires can be determined in a method as described below. A MODE solver is used to determine an effective refractive index  $n_{eff}$  of the nanowire array at the wavelength of interest  $\lambda_{air}$ , given the pitch of the nanowire array and the radius and material of the fluorescent nanowires therein. An effective wavelength  $\lambda_{eff}$  is defined as  $\lambda_{eff} = \lambda_{air} / n_{eff}$ . The length of the fluorescent nanowires is selected to be a multiple of  $\lambda_{eff}$ , which guarantees that the nanowire array has a collective mode at the wavelength of interest.

A MODE solver can accurately describe light propagation in a structure of any geometries. FDTD and MODE solvers from Lumerical Solutions, Inc. located at Vancouver, British Columbia, Canada were used.

The collective mode of the nanowire array can be an air mode. As shown in FIG. 7, the term “air mode” as used herein means a mode in which light intensity **700** is substantially zero within the fluorescent nanowires **120** and substantially non-zero outside the fluorescent nanowires **120**.

FIG. 8 shows a simulated fluorescence spectrum of an exemplary nanowire array **100** in axial directions thereof, using the FDTD method. The fluorescent nanowires **120** of this exemplary nanowire array **100** has a pitch of 637 nm and a radius of 60 nm. The fluorescent nanowires **120** of this exemplary nanowire array **100** are embedded in a material **150** which is an oxide. Compared with the spectra of FIG. 6, it can be seen that embedding the fluorescent nanowires **120** in the material **150** shifts the radius that maximizes fluorescence intensity of the nanowire array **100** from about 55 nm to about 60 nm. A partially reflective layer **180** can be deposited in areas on the surface **150a** around the fluorescent nanowires **120** as shown in FIGS. 2 and 3. The partially reflective layer **180** and the concave shape of the surface **150a** can enhance sharpness of the peak **800**. Sharpness of a peak can be defined by the Q value (Q factor, or quality factor) defined as center of

the peak divided by a full width at half maximum (FWHM) of the peak. A higher Q value indicates a sharper peak.

FIG. 9 shows a simulated fluorescence spectrum of an exemplary nanowire array **100** in axial directions thereof, using the FDTD method. The fluorescent nanowires **120** of this exemplary nanowire array **100** has a pitch of 1 micron, a radius of 55 nm and a height of 3 microns. The fluorescent nanowires **120** have a cladding layer **160** having a thickness of 300 nm. The fluorescent nanowires **120** are not embedded in a material **150**. The effect of the cladding layer **160** is evident by comparing the spectrum of FIG. 9 with the spectrum of FIG. 8: the peak **900** of FIG. 9 is taller than the peak **800** of FIG. 8 and the peak **900** is shifted to 750 nm from the position of the peak **800** at 637 nm.

The peak **900** can be adjusted to 637 nm by scaling the physical dimensions of the nanowire array **100** (pitch, nanowire radius, nanowire length) by a factor of 637 nm/750 nm=0.85. FIG. 10 shows result of this scaling. FIG. 10 shows a simulated fluorescence spectrum of an exemplary nanowire array **100** in axial directions thereof, using the FDTD method. The fluorescent nanowires **120** of this exemplary nanowire array **100** has a pitch of 0.85 micron (=1 micron $\times$ 0.85), a radius of 47 nm (=55 nm $\times$ 0.85) and a height of 2.55 microns (=3 microns $\times$ 0.85). The fluorescent nanowires **120** have a cladding layer **160** having a thickness of 255 nm (=300 nm $\times$ 0.85). Indeed the peak **1000** of FIG. 10 falls at 637 nm.

The nanowire array **100** can be used as a light source such as a laser. According to an embodiment, a light source comprises one or more of the nanowire array **100**. A method of using the nanowire array **100** as a light source comprises exposing the nanowire array **100** to a light, wherein the light preferable has a shorter wavelength than the wavelength of the wavelength of the collective mode of the nanowire array at which the fluorescent nanowires are operable to fluoresce. For example, if the wavelength of the collective mode is 500 nm, the light preferably has a wavelength less than 500 nm. The light can be provided by any suitable source such as LEDs, fluorescent lamps, mercury-vapor lamps, sodium-vapor lamps, discharge lamps, sunlight, incandescent lamps and/or laser.

The nanowire array **100** can be manufactured using an exemplary method, according to an embodiment, the exemplary method comprising: providing a substrate **110** having a layer of fluorescent material; coating a resist layer on the layer of fluorescent material; patterning the resist layer using a lithographic technique such as photolithography, e-beam lithography or holographic lithography such that a pattern corresponding to the fluorescent nanowires **120** is formed in the resist layer; forming the fluorescent nanowires **120** by etching the layer of fluorescent material; depositing the reflective layer **140** using a suitable deposition technique such as e-beam evaporation, thermal evaporation, or sputtering.

Another exemplary method of manufacturing the nanowire array **100**, according to an embodiment, comprises: providing a substrate **110**; growing the fluorescent nanowires **120** on the substrate by a suitable method such as the vapor-liquid-solid (VLS) method; depositing the reflective layer **140** using a suitable deposition technique such as e-beam evaporation, thermal evaporation, or sputtering. The VLS method is a method for the growth of one-dimensional structures, such as nanowires, from chemical vapor deposition. Growth of a crystal through direct adsorption of a gas phase on to a solid surface is generally very slow. The VLS method circumvents this by introducing a catalytic liquid alloy phase which can rapidly adsorb a vapor to supersaturation levels, and from which crystal growth can subsequently occur from nucleated seeds at the liquid-solid interface. The physical characteris-

tics of nanowires grown in this manner depend, in a controllable way, upon the size and physical properties of the liquid alloy.

The exemplary methods can also comprise a step of depositing a fluorescent material on the fluorescent nanowires **120** by a suitable technique such as e-beam evaporation, thermal evaporation, sputtering, chemical vapor deposition (CVD), atomic layer deposition (ALD), electrochemical deposition, followed by an optional annealing step.

The exemplary methods can also comprise a step of embedding a fluorescent material in the fluorescent nanowires by a suitable technique such as ion implantation, irradiation with electrons, followed by an optional annealing step.

The exemplary methods can also comprise a step of coating a dielectric nanowire around a fluorescent or an active emitting material such a nanowire laser. By a suitable technique such as ALD or CVD.

The exemplary methods can also comprise a step of coating a cladding layer surrounding a sidewall **120b** of the fluorescent nanowires, using a suitable technique such as e-beam evaporation, thermal evaporation, sputtering, chemical vapor deposition (CVD), atomic layer deposition (ALD), electrochemical deposition.

The exemplary methods can also comprise a step of filling space between the fluorescent nanowires with the material **150** by a suitable technique such as e-beam evaporation, thermal evaporation, sputtering, chemical vapor deposition (CVD), atomic layer deposition (ALD), electrochemical deposition, followed by an optional step of planarization such as chemical-mechanical planarization (CMP).

The foregoing detailed description has set forth various embodiments of the devices and/or processes by the use of diagrams, flowcharts, and/or examples. Insofar as such diagrams, flowcharts, and/or examples contain one or more functions and/or operations, it will be understood by those within the art that each function and/or operation within such diagrams, flowcharts, or examples can be implemented, individually and/or collectively, by a wide range of hardware, software, firmware, or virtually any combination thereof.

Those skilled in the art will recognize that it is common within the art to describe devices and/or processes in the fashion set forth herein, and thereafter use engineering practices to integrate such described devices and/or processes into data processing systems. That is, at least a portion of the devices and/or processes described herein can be integrated into a data processing system via a reasonable amount of experimentation.

The subject matter described herein sometimes illustrates different components contained within, or connected with, other components. It is to be understood that such depicted architectures are merely exemplary, and that in fact many other architectures can be implemented which achieve the same functionality. In a conceptual sense, any arrangement of components to achieve the same functionality is effectively "associated" such that the desired functionality is achieved. Hence, any two components herein combined to achieve a particular functionality can be seen as "associated with" each other such that the desired functionality is achieved, irrespective of architectures or intermediate components.

With respect to the use of substantially any plural and/or singular terms herein, those having skill in the art can translate from the plural to the singular and/or from the singular to the plural as is appropriate to the context and/or application. The various singular/plural permutations may be expressly set forth herein for sake of clarity.



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All references, including but not limited to patents, patent applications, and non-patent literature are hereby incorporated by reference herein in their entirety.

While various aspects and embodiments have been disclosed herein, other aspects and embodiments will be apparent to those skilled in the art. The various aspects and embodiments disclosed herein are for purposes of illustration and are not intended to be limiting, with the true scope and spirit being indicated by the following claims.

What is claimed is:

1. A nanowire array, comprising a substrate, a plurality of nanowires extending essentially perpendicularly from the substrate, wherein the nanowire array has a collective mode, wherein the nanowire array is configured to fluoresce at a wavelength of the collective mode.

2. The nanowire array of claim 1, further comprising a reflective layer at least disposed on the substrate in at least a portion of areas between the nanowires.

3. The nanowire array of claim 1, wherein the nanowires comprise fluorescent nanowires.

4. The nanowire array of claim 1, wherein the substrate comprises a material comprising silicon, sapphire, glass, silicon oxide, diamond, ZnO, a transparent non-absorbing dielectric material, or a combination thereof; or wherein the substrate has a thickness of about 0.1 to about 1 mm; or wherein the substrate comprises substantially a same material as or different materials from the nanowires.

5. The nanowire array of claim 2, wherein the reflective layer comprises a material selected from a group consisting of ZnO, Al, Au, Ag, Pd, Cr, Cu, and a combination thereof; or wherein the reflective layer has a reflectance of at least 50% for fluoresced light from the fluorescent nanowires; or wherein the reflective layer has a thickness of at least about 5 nm.

6. The nanowire array of claim 2, wherein the reflective layer is also disposed on the substrate in at least a portion of areas covered by the nanowires.

7. The nanowire array of claim 1, wherein the nanowires are at least partially embedded in a material.

8. The nanowire array of claim 1, wherein the nanowires comprise one or more fluorescent material.

9. The nanowire array of claim 1, wherein the nanowires comprise a material selected from the group consisting of inorganic fluorescent materials, organic fluorescent materials, quantum dots, diamond with color centers embedded therein, and a combination thereof.

10. The nanowire array of claim 1, wherein the nanowires comprise one or more non-fluorescent material with one or more fluorescent material embedded therein or covered thereby.

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11. The nanowire array of claim 1, wherein fluoresced light of the nanowire array is substantially spatially and/or temporally coherent.

12. The nanowire array of claim 1, wherein the wavelength of the collective mode of the nanowire array is substantially equal to a pitch of the nanowire array.

13. The nanowire array of claim 1, wherein the collective mode is an air mode.

14. A light source comprising one or more of the nanowire array of claim 1.

15. A method of using a nanowire array comprising exposing the nanowire array to light, wherein the nanowire array comprises:

a substrate,

a plurality of nanowires extending essentially perpendicularly from the substrate,

wherein the nanowire array has a collective mode, wherein the nanowire array is configured to fluoresce at a wavelength of the collective mode.

16. The method of claim 15, wherein the light has a shorter wavelength than the wavelength of the wavelength of the collective mode of the nanowire array.

17. The method of claim 15, wherein the light is provided by a source selected from the group consisting of LEDs, fluorescent lamps, mercury-vapor lamps, sodium-vapor lamps, discharge lamps, sunlight, incandescent lamps, laser and combinations thereof.

18. A method of manufacturing a nanowire array comprising:

obtaining a substrate;

forming a plurality of nanowires extend essentially perpendicularly from the substrate;

wherein the nanowire array is configured to fluoresce at a wavelength of a collective mode of the nanowire array.

19. The method of claim 18, wherein the substrate comprises a layer of fluorescent material, wherein forming the plurality of nanowires comprises etching the layer of fluorescent material.

20. The method of claim 18, wherein forming the plurality of nanowires comprises:

depositing a fluorescent material on or embedding a fluorescent material in the nanowires.

21. The method of claim 20, further comprising annealing the nanowires.

22. The method of claim 18, further comprising depositing a reflective layer at least on the substrate in at least a portion of areas between the nanowires.

23. The light source of claim 14, wherein the light source is a laser.

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